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NEWS 1 Web Page URLs for STN Seminar Schedule - N. America  
NEWS 2 Apr 08 "Ask CAS" for self-help around the clock  
NEWS 3 Apr 09 BEILSTEIN: Reload and Implementation of a New Subject Area  
NEWS 4 Apr 09 ZDB will be removed from STN

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NEWS 5 Apr 19 US Patent Applications available in IFICDB, IFIPAT, and IFIUDB  
NEWS 6 Apr 22 Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS

NEWS 7 Apr 22 BIOSIS Gene Names now available in TOXCENTER  
NEWS 8 Apr 22 Federal Research in Progress (FEDRIP) now available  
NEWS 9 Jun 03 New e-mail delivery for search results now available  
NEWS 10 Jun 10 MEDLINE Reload  
NEWS 11 Jun 10 PCTFULL has been reloaded  
NEWS 12 Jul 02 FOREGE no longer contains STANDARDS file segment  
NEWS 13 Jul 22 USAN to be reloaded July 28, 2002;  
saved answer sets no longer valid  
NEWS 14 Jul 29 Enhanced polymer searching in REGISTRY  
NEWS 15 Jul 30 NETFIRST to be removed from STN  
NEWS 16 Aug 08 CANCERLIT reload  
NEWS 17 Aug 08 PHARMAMarketLetter(PHARMAML) - new on STN  
NEWS 18 Aug 08 NTIS has been reloaded and enhanced  
NEWS 19 Aug 19 Aquatic Toxicity Information Retrieval (AQUIRE)  
now available on STN  
NEWS 20 Aug 19 IFIPAT, IFICDB, and IFIUDB have been reloaded  
NEWS 21 Aug 19 The MEDLINE file segment of TOXCENTER has been reloaded  
NEWS 22 Aug 26 Sequence searching in REGISTRY enhanced  
NEWS 23 Sep 03 JAPIO has been reloaded and enhanced  
NEWS 24 Sep 16 Experimental properties added to the REGISTRY file  
NEWS 25 Sep 16 Indexing added to some pre-1967 records in CA/CAPLUS  
NEWS 26 Sep 16 CA Section Thesaurus available in CAPLUS and CA  
NEWS 27 Oct 01 CASREACT Enriched with Reactions from 1907 to 1985

NEWS EXPRESS October 14 CURRENT WINDOWS VERSION IS V6.01,  
CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP),  
AND CURRENT DISCOVER FILE IS DATED 01 OCTOBER 2002  
NEWS HOURS STN Operating Hours Plus Help Desk Availability  
NEWS INTER General Internet Information  
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NEWS WWW      CAS World Wide Web Site (general information)

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FILE 'HOME' ENTERED AT 16:49:30 ON 17 OCT 2003

=> file registry

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 16:49:42 ON 17 OCT 2002  
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STRUCTURE FILE UPDATES: 16 OCT 2002 HIGHEST RN 462058-01-1  
DICTIONARY FILE UPDATES: 16 OCT 2002 HIGHEST RN 462058-01-1

TSCA INFORMATION NOW CURRENT THROUGH APRIL 2013

Please note that search-term pricing does apply when conducting a search.

Crossover limits based on the same approach as the one used by [Korpi et al. \(2010\)](#).

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:  
<http://www.cas.org/ONLINE/STN/STNOTES/stnote27.pdf>

1

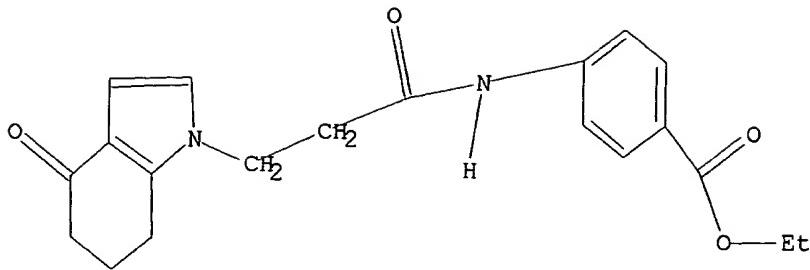
Uploading 29839388.htm

#### L1 STRUCTURE UPLOADED

⇒ d. 11

L1 HAS NO ANSWERS

BT HAS NO ANSWERS



Structure attributes must be viewed using STN Express query preparation.

=> s 11

SAMPLE SEARCH INITIATED 16:50:08 FILE 'REGISTRY'

~~SAMPLE SCREEN SEARCH COMPLETED - 3 TO ITERATE~~

100.0% PROCESSED            3 ITERATIONS  
SEARCH TIME: 00.00.01

1 ANSWERS

FULL FILE PROJECTIONS:    ONLINE    \*\*COMPLETE\*\*  
                              BATCH    \*\*COMPLETE\*\*

PROJECTED ITERATIONS:     3 TO      163

PROJECTED ANSWERS:        1 TO      80

L2                          1 SEA SSS SAM L1

=> s 11 sss full

FULL SEARCH INITIATED 16:50:16 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 12 TO ITERATE

100.0% PROCESSED            12 ITERATIONS  
SEARCH TIME: 00.00.02

1 ANSWERS

L3                          1 SEA SSS FUL L1

=> file caplus

COST IN U.S. DOLLARS

FULL ESTIMATED COST	SINCE FILE ENTRY	TOTAL SESSION
	140.28	140.49

FILE 'CAPLUS' ENTERED AT 16:50:24 ON 17 OCT 2002  
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FILE COVERS 1907 - 17 Oct 2002 VOL 137 ISS 16  
FILE LAST UPDATED: 16 Oct 2002 (20021016/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

=> s 11

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**REGISTRY INITIATED**

Substance data SEARCH and crossover from CAS REGISTRY in progress...  
Use DISPLAY HITSTR (or FHITSTR) to directly view retrieved structures.

SAMPLE SEARCH INITIATED 16:50:36 FILE 'REGISTRY'  
SAMPLE SCREEN SEARCH COMPLETED -            3 TO ITERATE

100.0% PROCESSED            3 ITERATIONS                            1 ANSWERS  
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:    ONLINE    \*\*COMPLETE\*\*  
                              BATCH     \*\*COMPLETE\*\*  
PROJECTED ITERATIONS:      3 TO        163  
PROJECTED ANSWERS:          1 TO        80

L4                            1 SEA SSS SAM L1

L5                            5 L4

=> d f bib hitstr abs total  
'F' IS NOT A VALID FORMAT FOR FILE 'CAPLUS'

The following are valid formats:

ABS ----- GI and AB  
ALL ----- BIB, AB, IND, RE  
APPS ----- AI, PRAI  
BIB ----- AN, plus Bibliographic Data and PI table (default)  
CAN ----- List of CA abstract numbers without answer numbers  
CBIB ----- AN, plus Compressed Bibliographic Data

DALL ----- ALL, delimited (end of each field identified)  
DMAX ----- MAX, delimited for post-processing  
FAM ----- AN, PI and PRAI in table, plus Patent Family data  
FBIB ----- AN, BIB, plus Patent FAM  
IND ----- Indexing data  
IPC ----- International Patent Classifications  
MAX ----- ALL, plus Patent FAM, RE  
PATS ----- PI, SO  
SAM ----- CC, SX, TI, ST, IT  
SCAN ----- CC, SX, TI, ST, IT (random display, no answer numbers;  
          SCAN must be entered on the same line as the DISPLAY,  
          e.g., D SCAN or DISPLAY SCAN)  
STD ----- BIB, IPC, and NCL

---

IABS ----- ABS, indented with text labels  
IALL ----- ALL, indented with text labels  
IBIB ----- BIB, indented with text labels  
IMAX ----- MAX, indented with text labels  
ISTD ----- STD, indented with text labels

OBIB ----- AN, plus Bibliographic Data (original)  
OIBIB ----- OBIB, indented with text labels

SBIB ----- BIB, no citations  
SIBIB ----- IBIB, no citations

HIT ----- Fields containing hit terms  
HITIND ----- IC, ICA, ICI, NCL, CC and index field (ST and IT)  
              containing hit terms  
HITRN ----- HIT RN and its text modification  
HITSTR ----- HIT RN, its text modification, its CA index name, and  
              its structure diagram  
HITSEQ ----- HIT RN, its text modification, its CA index name, its  
              structure diagram, plus NTE and SEQ fields  
FHITSTR ----- First HIT RN, its text modification, its CA index name, and  
              its structure diagram  
FHITSEQ ----- First HIT RN, its text modification, its CA index name, its  
              structure diagram, plus NTE and SEQ fields  
KWIC ----- Hit term plus 20 words on either side  
OCC ----- Number of occurrence of hit term and field in which it occurs

To display a particular field or fields, enter the display field codes. For a list of the display field codes, enter HELP DFIELDS at an arrow prompt (>). Examples of formats include: TI; TI,AU; BIB,ST; TI,IND; TI,SO. You may specify the format fields in any order and the information will be displayed in the same order as the format specification.

All of the formats (except for SAM, SCAN, HIT, HITIND, HITRN, HITSTR, FHITSTR, HITSEQ, FHITSEQ, KWIC, and OCC) may be used with DISPLAY ACC to view a specified Accession Number.

ENTER DISPLAY FORMAT (BIB):bib

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS

AN 2002:51464 CAPLUS  
 DN 136:112673  
 TI Methods using a purine derivative, pyrimidine derivative, or tetrahydroindolone derivative for treatment of disease-induced peripheral neuropathy and related conditions  
 IN Diamond, Jack; Glasky, Alvin J.  
 PA Neotherapeutics, Inc., USA  
 SO PCT Int. Appl., 69 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004452	A2	20020117	WO 2001-US21526	20010706
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
	AU 2001071908	A5	20020121	AU 2001-71908	20010706
	US 2002055506	A1	20020509	US 2001-900844	20010706
	US 2002061899	A1	20020523	US 2001-899901	20010706
PRAI	US 2000-216844P	P	20000707		
	WO 2001-US21526	W	20010706		
OS	MARPAT 136:112673				

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2002 ACS  
 AN 2002:51463 CAPLUS  
 DN 136:112672  
 TI Methods using a purine derivative, pyrimidine derivative, or tetrahydroindolone derivative for stimulation of synthesis of synaptophysin in the central nervous system  
 IN Glasky, Michelle; Lahiri, Debomoy K.; Farlow, Martin R.  
 PA Neotherapeutics, Inc., USA  
 SO PCT Int. Appl., 59 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004451	A2	20020117	WO 2001-US21385	20010706
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
	RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,				

DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,  
 BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG  
 US 2002040032 A1 20020404 US 2001-899478 20010705  
 AU 2001073218 A5 20020121 AU 2001-73218 20010706  
 PRAI US 2000-216808P P 20000707  
 WO 2001-US21385 W 20010706  
 OS MARPAT 136:112672

L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2002 ACS  
 AN 2002:51462 CAPLUS  
 DN 136:112671  
 TI Methods using a purine derivative, pyrimidine derivative, or tetrahydroindolone derivative for prevention of accumulation of amyloid .beta. peptide in the central nervous system  
 IN Glasky, Michelle; Lahiri, Debomoy K.; Farlow, Martin R.  
 PA Neotherapeutics, Inc., USA  
 SO PCT Int. Appl., 56 pp.  
 CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004450	A2	20020117	WO 2001-US21384	20010706
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	AU 2001073217	A5	20020121	AU 2001-73217	20010706
PRAI	US 2000-216845P	P	20000707		
	WO 2001-US21384	W	20010706		
OS	MARPAT	136:112671			

L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2002 ACS

AN 2002:51461 CAPLUS

DN 136:112691

TI Methods using a purine derivative, a pyrimidine derivative or a tetrahydroindolone derivative for treatment of conditions affected by activity of multidrug transporters

IN Taylor, Eve M.

PA Neotherapeutics, Inc., USA

SO PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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PI WO 2002004449 A2 20020117 WO 2001-US21383 20010706  
 WO 2002004449 A3 20020613

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,  
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 LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU,  
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 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY,  
 DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,  
 BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

US 2002128264 A1 20020912 US 2001-900297 20010706  
 PRAI US 2000-216616P P 20000707  
 OS MARPAT 136:112691

L5 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2002 ACS  
 AN 2002:51460 CAPLUS

DN 136:112670

TI Methods using purine derivatives, pyrimidine derivatives, and tetrahydroindolone derivatives for treatment of drug-induced peripheral neuropathy and related conditions

IN Diamond, Jack; Glasky, Alvin J.

PA Neotherapeutics, Inc., USA

SO PCT Int. Appl., 66 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004448	A2	20020117	WO 2001-US21373	20010706
	W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
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	US 2002061899	A1	20020523	US 2001-899901	20010706
PRAI	US 2000-216844P	P	20000707		
	WO 2001-US21373	W	20010706		
OS	MARPAT	136:112670			

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COST IN U.S. DOLLARS

FULL ESTIMATED COST

SINCE FILE ENTRY	TOTAL SESSION
5.57	146.84

STN INTERNATIONAL LOGOFF AT 16:51:42 ON 17 OCT 2002

CODEN: PIXXD2

DT Patent  
 LA English  
 FAN.CNT 2

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004452	A2	20020117	WO 2001-US21526	20010706
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AU	2001071908	A5	20020121	AU 2001-71908	20010706
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US	2002055506	A1	20020509	WO 2001-US21526W	20010706
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US	2002061899	A1	20020523	US 2000-216844PP	20000707
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				US 2000-216844PP	20000707

## PATENT FAMILY INFORMATION:

FAN 2002:51460

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004448	A2	20020117	WO 2001-US21373	20010706
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AU	2001073212	A5	20020121	AU 2001-73212	20010706
				US 2000-216844PP	20000707
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				US 2001-900844	20010706
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				US 2000-216844PP	20000707

OS MARPAT 136:112673

IT 389799-42-2

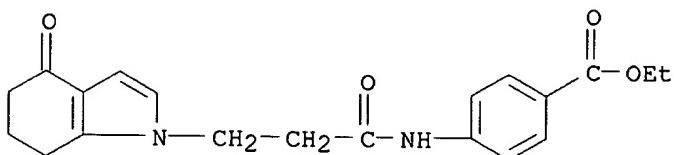
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
 (purine derivs., pyrimidine derivs., and tetrahydroindolone derivs.)

for

treatment of disease-induced peripheral neuropathy and related conditions)

RN 389799-42-2 CAPLUS

CN Benzoic acid, 4-[[1-oxo-3-(4,5,6,7-tetrahydro-4-oxo-1H-indol-1-yl)propyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



**AB** A method of treating disease-induced peripheral neuropathy comprises administering to a patient with disease-induced peripheral neuropathy an effective quantity of a purine deriv. or analog, a tetrahydroindolone deriv. or analog, or a pyrimidine deriv. or analog. If the compd. is a purine deriv., the purine moiety can be guanine or hypoxanthine. The compd. can induce peripheral nerve sprouting through the action of a neurotrophic factor such as nerve growth factor (NGF) without the occurrence of hyperalgesia. The peripheral nerve sprouting can be nociceptive nerve sprouting. The disease-induced peripheral neuropathy can be diabetic neuropathy or disease-induced peripheral neuropathy with another basis.

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2002 ACS

AN 2002:51463 CAPLUS

DN 136:112672

TI Methods using a purine derivative, pyrimidine derivative, or tetrahydroindolone derivative for stimulation of synthesis of synaptophysin in the central nervous system

IN Glasky, Michelle; Lahiri, Debomoy K.; Farlow, Martin R.

PA Neotherapeutics, Inc., USA

SO PCT Int. Appl., 59 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004451	A2	20020117	WO 2001-US21385	20010706
	W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG		US 2000-216808PP	20000707
US	2002040032	A1	20020404	US 2001-899478	20010705
				US 2000-216808PP	20000707
AU	2001073218	A5	20020121	AU 2001-73218	20010706

US 2000-216808PP 20000707  
WO 2001-US21385W 20010706

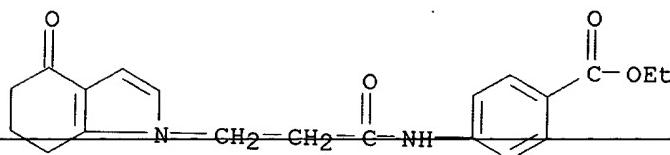
OS MARPAT 136:112672

IT 389799-42-2

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
(purine deriv., pyrimidine deriv., or tetrahydroindolone deriv. for stimulation of synthesis of synaptophysin in CNS)

RN 389799-42-2 CAPLUS

CN Benzoic acid, 4-[[1-oxo-3-(4,5,6,7-tetrahydro-4-oxo-1H-indol-1-yl)propyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



AB A method of increasing the synthesis and/or secretion of synaptophysin comprises administering to a patient with a neurolog. disease or a patient at risk of developing a neurolog. disease an effective quantity of a purine deriv. of analog, a tetrahydroindolone deriv. or analog, or a pyrimidine deriv. or analog. If the compd. is a purine deriv., the purine moiety can be guanine or hypoxanthine. The neurolog. disease can be a neurodegenerative disease such as Alzheimer's disease or a neurodevelopmental disorder such as Down's syndrome. Typically, the compd. can pass through the blood-brain barrier. A particularly preferred purine deriv. is N-4-carboxyphenyl-3-(6-oxohypoxanthine-9-yl)propanamide.

LS ANSWER 3 OF 5 CAPLUS COPYRIGHT 2002 ACS

AN 2002:51462 CAPLUS

DN 136:112671

TI Methods using a purine derivative, pyrimidine derivative, or tetrahydroindolone derivative for prevention of accumulation of amyloid .beta. peptide in the central nervous system

IN Glasky, Michelle; Lahiri, Debomoy K.; Farlow, Martin R.

PA Neotherapeutics, Inc., USA

SO PCT Int. Appl., 56 pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE	
PI	WO 2002004450	A2	20020117	WO 2001-US21384	20010706	
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 BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG

US 2002040031 A1 20020404 US 2000-216845PP 20000707  
 AU 2001073217 A5 20020121 US 2001-899611 20010705  
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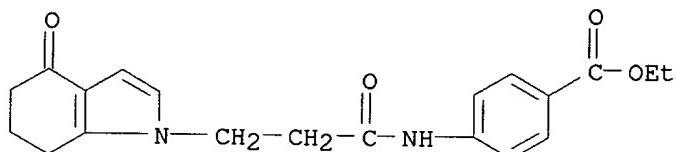
OS MARPAT 136:112671

IT 389799-42-2

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)  
 (purine deriv., pyrimidine deriv., or tetrahydroindolone deriv. for prevention of accumulation of amyloid .beta. peptide in CNS)

RN 389799-42-2 CAPLUS

CN Benzoic acid, 4-[[1-oxo-3-(4,5,6,7-tetrahydro-4-oxo-1H-indol-1-yl)propyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



AB A method of either inhibiting the formation of A.beta. or stimulating the formation of sAPP comprises administering to a patient with a neurol. disease or a patient at risk of developing a neurol. disease an effective quantity of a purine deriv. or analog, a tetrahydroindolone deriv. or analog, or a pyrimidine deriv. or analog. If the compd. is a purine deriv., the purine moiety can be guanine or hypoxanthine. The neurol. disease can be a neurodegenerative disease such as Alzheimer's disease or a neurodevelopmental disorder such as Down's syndrome. Typically, the compd. can pass through the blood-brain barrier. A particularly preferred purine deriv. is N-4- carboxyphenyl-3-(6-oxohydropurin-9-yl)propanamide.

L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2002 ACS  
 AN 2002:51461 CAPLUS

DN 136:112691

TI Methods using a purine derivative, a pyrimidine derivative or a tetrahydroindolone derivative for treatment of conditions affected by activity of multidrug transporters

IN Taylor, Eve M.

PA Neotherapeutics, Inc., USA

SO PCT Int. Appl., 70 pp.

CODEN: PIXXD2

DT Patent

LA English

## FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004449	A2	20020117	WO 2001-US21383	20010706
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OS MARPAT 136:112691

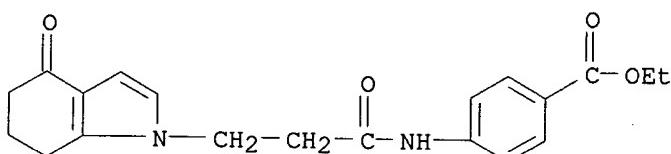
~~IT 389799-42-2~~

RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(purine deriv., pyrimidine deriv. or tetrahydroindolone deriv. for treatment of conditions affected by activity of multidrug transporters)

RN 389799-42-2 CAPLUS

CN Benzoic acid, 4-[[1-oxo-3-(4,5,6,7-tetrahydro-4-oxo-1H-indol-1-yl)propyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



AB One aspect of the invention is a method of treating a condition or disease

assocd. with the activity of a multidrug transporter protein comprising administering to a mammal with a condition or disease assocd. with the activity of a multidrug transporter protein an effective quantity of a purine deriv. or analog, a tetrahydroindolone deriv. or analog, or a pyrimidine deriv. or analog. If the compd. is a purine deriv., the purine

moiety can be guanine or hypoxanthine. A particularly preferred bifunctional purine deriv. is N-4-carboxyphenyl-3-(6-oxohypoxanthine-9-yl)propanamide. The methods of the invention can be used to treat cancer,

a microbial or parasitic infection, HIV, infection, or a condition assocd.

with inflammation, e.g. asthma or rheumatic disease.

L5 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2002 ACS

AN 2002:51460 CAPLUS  
 DN 136:112670  
 TI Methods using purine derivatives, pyrimidine derivatives, and tetrahydroindolone derivatives for treatment of drug-induced peripheral neuropathy and related conditions  
 IN Diamond, Jack; Glasky, Alvin J.  
 PA Neotherapeutics, Inc., USA  
 SO PCT Int. Appl., 66 pp.  
 CODEN: PIXXD2  
 DT Patent  
 LA English  
 FAN.CNT 2

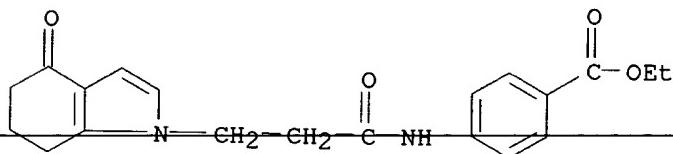
	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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	AU 2001073212	A5	20020121	AU 2001-73212	20010706
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	US 2002055506	A1	20020509	US 2001-900844	20010706
				US 2000-216844PP	20000707
	US 2002061899	A1	20020523	US 2001-899901	20010706
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## PATENT FAMILY INFORMATION:

FAN 2002:51464

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2002004452	A2	20020117	WO 2001-US21526	20010706
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				WO 2001-US21526W	20010706
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	US 2002061899	A1	20020523	US 2001-899901	20010706
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OS MARPAT 136:112670  
 IT 389799-42-2  
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL  
 (Biological study); USES (Uses)  
 (purine derivs., pyrimidine derivs., and tetrahydroindolone derivs.  
 for treatment of drug-induced peripheral neuropathy and related  
 conditions)  
 RN 389799-42-2 CAPLUS  
 CN Benzoic acid, 4-[[1-oxo-3-(4,5,6,7-tetrahydro-4-oxo-1H-indol-1-yl)propyl]amino]-, ethyl ester (9CI) (CA INDEX NAME)



AB A method of treating drug-induced peripheral neuropathy comprises administering to a patient with drug-induced peripheral neuropathy an effective quantity of a purine deriv. or analog, a tetrahydroindolone deriv. or analog, or a pyrimidine deriv. or analog. If the compd. is a purine deriv., the purine moiety can be guanine or hypoxanthine. The compd. can induce peripheral nerve sprouting through the action of a neurotrophic factor such as nerve growth factor (NGF) without the occurrence of hyperalgesia. The peripheral nerve sprouting can be nociceptive nerve sprouting. The drug-induced peripheral neuropathy can be drug-induced peripheral neuropathy assocd. with the administration of oncolytic drugs, such as a vinca alkaloid, cisplatin, paclitaxel, suramin, altretamine, carboplatin, chlorambucil, cytarabine, dacarbazine, docetaxel, etoposide, fludarabine, ifosfamide with mesna, tamoxifen, teniposide, or thioguanine. The methods of the invention are particularly useful in treating peripheral neuropathy assocd. with the administration of vincristine, paclitaxel, or cisplatin.

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DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
CA SUBSCRIBER PRICE	ENTRY	SESSION
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<10/17/2002>